Amendments to the Claims

This listing of the claims as amended will replace all prior versions, and listings, of claims in the application.

- 1-55. (Canceled)
- 56. (Currently amended) A method of treating a subject having a disorder or condition characterized by an unwanted immune response comprising administering an effective amount of an early activation molecule depletor a depleting anti-CD69 antibody molecule, wherein the anti-CD69 antibody specifically binds SEQ ID NO:2 to the subject.
 - 57-58. (Canceled)
- 59. (Currently amended) The method of claim 5856, wherein said depletor is a depleting anti-CD69 antibody molecule selected from the group consisting of: a humanized anti-CD69 antibody molecule, a human anti-CD69 antibody molecule, a chimeric anti-CD69 antibody molecule and a deimmunized anti-CD69 antibody molecule.
- 60. (Original) The method of claim 59, wherein said human anti-CD69 antibody molecule is a monoclonal antibody.
 - 61-66. (Canceled)
- 67. (Original) The method of claim 56, wherein said disorder is an acute or chronic inflammatory disorder, or an immune disorder.
- 68. (Original) The method of claim 67, wherein said disorder is an autoimmune disorder.
- 69. (Original) The method of claim 56, wherein the disorder is selected from the group consisting of: rheumatoid arthritis, systemic lupus erythematosus, scleroderma, Sjögren

syndrome, autoimmune diabetes, thyroiditis, and other organo-specific immune diseases, including psoriasis.

- 70. (Withdrawn) The method of claim 56, wherein the disorder is a neurological disorder, a gastrointestinal disorder, a cardiovascular disorder or a respiratory disorder.
- 71. (Withdrawn) The method of claim 70, wherein the disorder is a neurological disorder and the neurological disorder is selected from the group consisting of: multiple sclerosis, myasthenia gravis, and other neurological immune-mediated diseases.
- 72. (Withdrawn) The method of claim 70, wherein the disorder is a gastrointestinal disorder and the gastrointestinal disorder is selected from the group consisting of Crohn's disease, colitis, celiac disease, and hepatitis.
- 73. (Withdrawn) The method of claim 70, wherein the disorder is a respiratory disorder and the respiratory disorder is selected from the group consisting of: emphysema, and respiratory airways infections.
- 74. (Withdrawn) The method of claim 70, wherein the disorder is a cardiovascular disorder and the cardiovascular disorder is selected from the group consisting of: atherosclerosis, cardiomyopathy, rheumatic fever, endocarditis, vasculitis, and other immune-mediated diseases.
- 75. (Withdrawn) The method of claim 56, wherein the disorder is an allergic process or a hypersensitivity reaction (type I, II, III, and IV), including asthma, rhinitis, and other immune-mediated hypersensitivity reactions.
- 76. (Withdrawn) The method of claim 56, wherein the disorder is transplant or graft rejection.
- 77. (Withdrawn) The method of claim 56, wherein said disorder or condition is: acute lung injury, acute respiratory distress syndrome, asthma, bronchitis, cystic fibrosis,

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reperfusion injury, nephritis, pancreatitis, artery occlusion, stroke, transplantation, ultraviolet light induced injury, vasculitis, and sarcoidosis.

78-104. (Canceled)

- 105. (Currently amended) A method of treating a subject having a disorder or condition characterized by an unwanted immune response comprising administering an effective amount of an antibody specific for an early activation molecule a depleting anti-CD69 antibody molecule, wherein the anti-CD69 antibody specifically binds SEQ ID NO:2 to the subject, alone or conjugated to a second therapeutic agent.
- 106. (Original) The method of claim 105, wherein said second therapeutic agent is selected from the group consisting of: chemotherapeutic agents; radioisotopes; and cytotoxins.
- 107. (Original) The method of claim 105, wherein the antibody is a monoclonal antibody.
- 108. (Original) The method of claim 107, wherein the monoclonal antibody is a human antibody.